Model for the spin-dependent Seebeck coefficient of InSb in a magnetic field

Nicholas A. Pike^{*} and David Stroud

Department of Physics, The Ohio State University, Columbus, Ohio 49210, USA (Received 22 September 2014; revised manuscript received 29 October 2014; published 25 November 2014)

We develop a simple theory for the spin-dependent Seebeck effect in *n*-doped InSb in an external magnetic field. We consider spin-1/2 electrons in the conduction band of InSb with a temperature gradient parallel to the applied magnetic field. In the absence of spin-orbit interactions, a Boltzmann equation approach leads to a spin current parallel to the field and proportional to the temperature gradient. The calculated spin-dependent Seebeck coefficient oscillates as a function of magnetic field *B*; the peak positions are approximately periodic in 1/B. The oscillations arise when the Fermi energy crosses the bottom of a Landau band.

DOI: 10.1103/PhysRevB.90.174435

PACS number(s): 72.25.Dc, 72.20.Pa, 72.15.Gd

I. INTRODUCTION

It is well known that spin currents can be generated in many materials by the flow of heat current. The so-called spin Seebeck effect arises when this spin motion leads to the production of a voltage, usually by the inverse spin Hall effect (ISHE). A number of recent experiments have demonstrated the occurrence of a spin Seebeck effect in a variety of materials [1-4] and in both transverse and longitudinal experimental configurations [2–4]. The materials involved can be metallic ferromagnets, magnetic insulators, and even doped nonmagnetic semiconductors (such as Tedoped InSb) in a strong magnetic field. Several papers have discussed possible explanations for such behavior in a range of materials [4–11]. Another related effect, which also arises from interaction of spins with heat currents, is known as the spin-dependent Seebeck effect. This involves the production of a bulk spin current by an applied temperature gradient, but unlike the spin-Seebeck effect does not require an ISHE, which arises from spin-orbit interactions.

In this paper, we present a simple calculation for spin transport, and specifically for the spin-dependent Seebeck coefficient in *n*-doped InSb in the presence of a temperature gradient and an external magnetic field. Our model is based on the Boltzmann equation, but applied to the bands formed by the Landau levels in an *n*-type semiconductor when there is a strong magnetic field parallel to the temperature gradient. The model readily leads to spin-dependent transport. The results we obtain are relevant to the spin-dependent Seebeck effect in *n*-doped InSb, but do not directly apply to the experiments of Ref. [1] where the transverse spin Seebeck coefficient was measured. The coefficient we calculate, however, does show the characteristic oscillations with inverse magnetic field seen in Ref. [1], and we believe that if the spin-dependent Seebeck effect were explicitly measured in *n*-doped InSb, similar oscillations would be found. In the future, by including spin-orbit interactions, it may be possible to calculate the measured transverse spin Seebeck effect in InSb.

The band structures of InSb and other zinc-blende semiconductors have been extensively investigated, both theoretically and experimentally. Early theoretical studies by Kane [12], Dresselhaus [13], and Parmenter [14] explain the effects of symmetry on the conduction band electronic states. In other early studies, the effects of a magnetic field on the band structure of InSb were investigated by Roth *et al.* [15,16] and by Pidgeon *et al.* [17]. These theoretical and experimental studies led to a better understanding of the beats observed in Shubnikov-de Haas oscillations in III-V semiconductors [18]. Other experiments showed that the lowest conduction band state in InSb has the spherically symmetric Γ_6 symmetry [13,19], and that the effective mass of the conduction band electrons is only a small fraction of the free electron mass [19].

The remainder of the paper is organized as follows. In Sec. II, we briefly review the relevant macroscopic transport equations describing the coupled heat, electronic, and spin transport. In Sec. III, we present a theory for these transport coefficients based on a microscopic Hamiltonian combined with the Boltzmann equation. The Hamiltonian includes the Landau Hamiltonian for electrons in a magnetic field and the Zeeman interaction between the spins and the magnetic field. The Boltzmann equation is then linearized, and solved to yield the thermoelectric and spin-dependent Seebeck coefficients. In Sec. IV, we present numerical solutions of this model for the various transport coefficients as a function of magnetic field at a temperature T = 4.5 K. In Sec. V, we give a brief concluding discussion. An Appendix gives explicit expressions for the various Onsager coefficients.

II. MACROSCOPIC TRANSPORT EQUATIONS

We begin by writing down the appropriate macroscopic transport equations for the system of interest, which we visualize as a doped semiconductor such as *n*-InSb in a magnetic field **B** taken parallel to the *z* axis. In this case, there are three current densities to consider: the heat current density J_Q , and the charge current densities J_+ and J_- for spin-up and spin-down charge carriers.

These currents are expected to be linearly related to corresponding fields, which will be $-\nabla T$, and $-\nabla \mu_{\pm}$, where T is the temperature and the μ_{\pm} are the electrochemical potentials for spin up and spin down charge carriers. As noted by Brechet and Ansermet [20] (see also Bauer *et al.* [4] and Uchida *et al.* [21]), the linear Onsager relations between these currents and fields may be written

^{*}Pike.55@osu.edu

as

$$\begin{pmatrix} \mathbf{J}_{\mathbf{Q}} \\ \mathbf{J}_{+} \\ \mathbf{J}_{-} \end{pmatrix} = \begin{pmatrix} L_{\mathcal{Q}\mathcal{Q}} & L_{\mathcal{Q}+} & L_{\mathcal{Q}-} \\ L_{+\mathcal{Q}} & L_{++} & L_{+-} \\ L_{-\mathcal{Q}} & L_{-+} & L_{--} \end{pmatrix} \begin{pmatrix} -\nabla T \\ -\nabla \mu_{+} \\ -\nabla \mu_{-} \end{pmatrix}.$$
(1)

Instead of the currents \mathbf{J}_+ and \mathbf{J}_- , it may be more convenient to consider the charge current density $\mathbf{J}_e = \mathbf{J}_+ + \mathbf{J}_-$ and the spin current density $\mathbf{J}_S = (\hbar/2q)(\mathbf{J}_+ - \mathbf{J}_-)$ (where q = -eis the charge of the current carriers and we assume a spin $\hbar/2$ per carrier). Similarly, rather than the $-\nabla \mu_{\pm}$, it is more convenient, following Valet and Fert [22], to introduce the quantities μ_{av} and $\Delta \mu$ via the relation

$$\mu_{\pm} = \mu_{av} \pm \Delta \mu + qV. \tag{2}$$

Here μ_{av} is the average of the two chemical potentials at zero applied voltage, V is the electrostatic potential, and $\Delta \mu = \frac{1}{2}(\mu_+ - \mu_-)$.

We can now write down the linear transport equations in terms of these new fields and currents. The result is

$$\begin{pmatrix} \mathbf{J}_{\mathbf{Q}} \\ \mathbf{J}_{e} \\ \mathbf{J}_{S} \end{pmatrix} = \begin{pmatrix} L_{\mathcal{Q}\mathcal{Q}} & L_{\mathcal{Q}e} & L_{\mathcal{Q}S} \\ L_{e\mathcal{Q}} & L_{ee} & L_{eS} \\ L_{S\mathcal{Q}} & L_{Se} & L_{SS} \end{pmatrix} \begin{pmatrix} -\nabla T \\ \mathcal{E} \\ -\frac{\hbar}{2q} \nabla(\Delta \mu) \end{pmatrix}, \quad (3)$$

where the various *L* coefficients are all certain linear combinations of the coefficients in Eq. (1) and $\mathcal{E} = \mathbf{E} + \frac{\nabla \mu}{|e|}$ is the effective electric field.

Equation (3) applies if the spin polarization is parallel to the direction of the spin current. If the spin polarization is not parallel to the direction of spin current flow, then there are three spin current vectors, to be called $\mathbf{J}_{S,i}$ with i = x, y, and z, corresponding to current densities of the x, y, and zcomponents of electron spin. In this case, Eq. (3) should be replaced by a 5×5 matrix equation, corresponding to the five current densities \mathbf{J}_Q , \mathbf{J}_e , and the three $\mathbf{J}_{S,i}$'s. Since we will not consider this situation in the present paper, we will not write down this equation explicitly.

III. THEORY FOR TRANSPORT COEFFICIENTS IN INSB

Next, we present a theory for some of the above transport coefficients in an *n*-type semiconductor, such as Te-doped InSb, in a magnetic field. We are particularly interested in the coefficient L_{SQ} , which is the induced spin current density in the *z* direction when a temperature gradient is applied in that direction, as indicated in Eq. (3). As discussed in Ref. [23], we envision that an experiment to measure this coefficient could be carried out in a wire that is part of an open electric circuit (no electrical current flowing), as shown schematically in Fig. 1. As indicated in the figure, both a uniform magnetic field **B** and a uniform temperature gradient ∇T are parallel to the wire, which is assumed to lie in the *z* direction. A further analysis of the relevant boundary conditions for this geometry is given later in the paper.

A. Electronic energies and wave functions

The conduction band of InSb is nondegenerate, and the low-lying electronic states in this band have the spectrum of a free electron (of effective mass m^*) in a magnetic field.



FIG. 1. (Color online) Schematic of an idealized experimental configuration, showing the orientation of the magnetic field, thermal gradient, and the positions of the nonmagnetic contacts (red) used to probe the spin-dependent Seebeck coefficient in InSb (shown in tan). The thermal gradient and applied magnetic field are oriented along *z*. Depending on the boundary conditions, as discussed this coefficient may be determined by the longitudinal voltage difference measured between the red contacts [see Eq. (24)] [4].

The spin-independent part of the effective-mass Hamiltonian describing these states is thus

$$H_{0} = \frac{1}{2m^{*}} \bigg[+ \bigg(-i\hbar \frac{\partial}{\partial x} + qBy \bigg)^{2} - \hbar^{2} \frac{\partial^{2}}{\partial y^{2}} - \hbar^{2} \frac{\partial^{2}}{\partial z^{2}} \bigg],$$
(4)

where we have used SI units and a gauge such that the vector potential $\mathbf{A} = (-By, 0, 0)$, where $\mathbf{B} = \nabla \times \mathbf{A}$ is the applied magnetic field.

The solutions of the spin-independent Hamiltonian given in Eq. (4) are standard. The total energy is a function of a wave vector k_z and the Landau level index n, and can be written as

$$E_n(k_z) = \frac{\hbar^2 k_z^2}{2m^*} + \left(n + \frac{1}{2}\right)\hbar\omega_c,$$
 (5)

where $\omega_c = eB/m^*$ is the cyclotron frequency. Each level has a degeneracy per spin σ_z of

$$N_{n,\sigma_z} = \frac{L_x L_y B}{\Phi_0},\tag{6}$$

where $\Phi_0 = h/2e$ is the magnetic flux quantum.

In the absence of spin-orbit interaction, the spin-dependent part of the electronic Hamiltonian, denoted H_s , consists of a Zeeman interaction between the conduction electron and the applied magnetic field, which may be written as

$$H_s = g\mu_B \mathbf{B} \cdot \boldsymbol{\sigma}. \tag{7}$$

Here g is the electronic g factor, which is assumed independent of the magnetic field strength, μ_B is the Bohr magneton, and σ is the vector of the three Pauli spin matrices for a spin-1/2 particle. The eigenvalues of $H_0 + H_s$ are characterized by quantum numbers n, k_z , and $\sigma_z = \pm 1/2$, and are given by

$$E_{n,\sigma_z}(k_z) = E_n(k_z) + g\mu_B B\sigma_z, \qquad (8)$$

with a spin σ_z parallel to z and a degeneracy given by Eq. (6).

B. Electrical, thermal, and spin currents

Next, we will obtain the various electronic transport coefficients for an *n*-type semiconductor such as InSb, using the Boltzmann equation. In the presence of a magnetic field, the conduction band is broken up into many one-dimensional bands, labeled by a Landau level index *n* and a spin index σ_z ($\sigma_z = \pm$). Each band is also highly degenerate, with degeneracy N_{n,σ_z} as given in Eq. (6). The Boltzmann equation

for an electron of spin σ_z in band *n* can be written in the standard way (see, e.g., Ref. [25]) as

$$\frac{\partial g_{n,\sigma_z}}{\partial t} + \mathbf{v}_{n\sigma_z}(k_z) \cdot \nabla_{\mathbf{r}} g_{n,\sigma_z} + \frac{\mathbf{F}}{\hbar} \cdot \nabla_{k_z} g_{n,\sigma_z} = \left(\frac{\partial g_{n,\sigma_z}}{\partial t}\right)_{\text{coll}}.$$
(9)

Here $g_{n,\sigma_z}(\mathbf{r},k_z,t)$ is the probability that an electron in a state k_z in the n^{th} band with spin σ_z at a position \mathbf{r} is occupied at time t, $\mathbf{F} = -e\mathcal{E} = -e\mathbf{E} - \nabla\mu$, and $\mathbf{v}_{n\sigma_z}(k_z)$ is the velocity of an electron in the state described by k_z , σ_z and n.

As is conventional, we make the relaxation time approximation so that the collision term is rewritten as

$$\left(\frac{\partial g_{n,\sigma_z}}{\partial t}\right)_{\text{coll}} \sim -\frac{\delta g_{n,\sigma_z}(\mathbf{r},k_z,t)}{\tau},\tag{10}$$

where $\delta g_{n,\sigma_z}$ is the deviation of g_{n,σ_z} from its equilibrium value g_{n,σ_z}^0 . The function g_{n,σ_z}^0 is set equal to the Fermi function given by

$$g_{n,\sigma_{z}}^{0}(k_{z}) = \frac{1}{\exp\left[\beta\left(E_{n,\sigma_{z}}(k_{z}) - \mu\right)\right] + 1},$$
 (11)

where $\beta = 1/k_B T$, T is the temperature, k_B is Boltzmann's constant, and μ is the chemical potential.

We seek a steady-state solution and thus the first term on the left-hand side of Eq. (9) vanishes. We also linearize the Boltzmann equation by assuming that both $\nabla_r g_{n,\sigma_z}$ and **F** are small, so that the factor g_{n,σ_z} in both the second and the third terms of Eq. (9) can be approximated as g_{n,σ_z}^0 . Combining these conditions, we obtain the linearized steady-state Boltzmann equation in the relaxation time approximation, which, after simplification, is

$$-\frac{\delta g_{n,\sigma_z}(k_z)}{\tau} = \frac{E_{n,\sigma_z}(k_z) - \mu}{T} \left(-\frac{\partial g_{n,\sigma_z}^0}{\partial E} \right) \nabla T \cdot \mathbf{v}_{n,\sigma_z}(k_z) + \mathbf{F} \cdot \mathbf{v}_{n,\sigma_z}(k_z) \left(\frac{\partial g_{n,\sigma_z}^0}{\partial E} \right), \tag{12}$$

where we evaluate *E* at $E = E_{n,\sigma_z}(k_z)$.

We are interested in the case of an applied temperature gradient and effective electric field oriented along the *z* axis [see Eq. (3)]. We can now express the electric, heat, and spin currents in terms of $\delta g_{n,\sigma_z}(k_z)$, as obtained from Eq. (12). Two of these expressions are given, for a spherical band, by, e.g., Ref. [25]. These general expressions need to be modified to take account of the degeneracy of the Landau bands as we do below. The mathematical form of the spin current density, \mathbf{J}_s , is similar to that of the electrical current density, \mathbf{J}_e .

To be explicit, we can write out the current densities in the system as follows:

$$\mathbf{J}_{e} = \frac{L_{z}}{V} \sum_{n,\sigma_{z}} \int \frac{dk_{z}}{2\pi} (-e) N_{n,\sigma_{z}} \mathbf{v}_{n,\sigma_{z}}(k_{z}) \delta g_{n,\sigma_{z}}(k_{z}), \qquad (13)$$

$$\mathbf{J}_{\mathcal{Q}} = \frac{L_z}{V} \sum_{n,\sigma_z} \int \frac{dk_z}{2\pi} N_{n,\sigma_z} \Delta E_{n,\sigma_z} \mathbf{v}_{n,\sigma_z}(k_z) \delta g_{n,\sigma_z}(k_z), \quad (14)$$

$$\mathbf{J}_{S,i} = \frac{L_z}{V} \sum_{n,\sigma_z} \int \frac{dk_z}{2\pi} \mu_B \langle \sigma_{n,i} \rangle N_{n,\sigma_z} \mathbf{v}_{n,\sigma_z}(k_z) \delta g_{n,\sigma_z}(k_z), \quad (15)$$

where the limits of the integral are $\pm \infty$ and $\Delta E_{n,\sigma_z} = E_{n,\sigma_z}(k_z) - \mu$.

In Eqs. (13)–(15), \mathbf{J}_Q is the heat current, $\mathbf{v}_{n,\sigma}(k_z) = \hbar^{-1} \nabla_k E_{n,\sigma_z}(k_z)$ is the velocity of the electron in the band labeled by (n,σ_z) , and $\langle \sigma_{n,i} \rangle$ is the expectation value of the *i*th component of spin in the band (n,σ_z) (i = x, y, z). In the absence of spin-orbit interaction, only $\mathbf{J}_{S,z}$, that is, the current density associated with the *z* component of spin, is nonzero.

For an electron in the conduction band the velocity, $\mathbf{v}_{n,\sigma_z}(k_z) = v_{n,\sigma_z}(k_z)\hat{z}$ is given by

$$v_{n,\sigma_z}(k_z) = \frac{\hbar k_z}{m^*}$$
$$= \pm \frac{\hbar}{m^*} \sqrt{\frac{2m^*}{\hbar^2} \left(E_{n,\sigma_z}(k_z) - E_{n,\sigma_z}^0 \right)}, \tag{16}$$

where the + and - signs apply when $k_z > 0$ and $k_z < 0$, respectively, and E_{n,σ_z}^0 is defined as the minimum energy for the band (n, σ_z) given by

$$E_{n,\sigma_z}^0 = \left(n + \frac{1}{2}\right)\hbar\omega_c + \sigma_z g\mu_B B.$$
(17)

In order to calculate the various transport coefficients in Eqs. (13)–(15), we need the chemical potential μ . μ can be calculated given the conduction electron density $\rho = N_e/V$, where N_e is the total number of conduction electrons in volume V. For the present case, the chemical potential μ is obtained from

$$\rho = \frac{B}{\pi \Phi_0} \sum_{n,\sigma_z} \int_0^\infty g_{n,\sigma_z}^0(k_z) dk_z \tag{18}$$

where Eq. (18) is an implicit equation for $\mu(T, B)$.

Since the experiments of Ref. [1] are done at a very low temperature (T = 4.5 K), we have approximated μ (or equivalently, the Fermi energy E_F) by its value at T = 0. In this case, g_{n,σ_z}^0 is just a step function, and E_F is given implicitly by

$$\rho = \sum_{n,\sigma_z} \frac{B}{\pi \Phi_0} \left(\frac{2m^*}{\hbar^2} (\mu - E_{n,\sigma_z}^0) \right)^{\frac{1}{2}},$$
 (19)

where E_{n,σ_z}^0 is defined in Eq. (17) and the sum runs only over Landau bands with nonzero electron occupation at T = 0.

C. Transport coefficients

We obtain the transport coefficients of interest by using Onsager's linear relationship between the currents and the forces generating the currents [26]. For the present problem, this relation is given by Eq. (3). These may be written in condensed form as

$$J_i = \sum_j L_{ij} F_j, \tag{20}$$

where i = e, Q, S runs over the three currents in the system and *j* runs over the forces acting on the conduction electrons. In this paper, we consider only longitudinal spin currents, i.e., we assume that only $\langle \sigma_z \rangle \neq 0$. The explicit form of the measured transport coefficients can then be obtained by combining Eq. (3) with Eqs. (12)–(16). For example, the electrical conductivity, σ_e , is given by the Onsager coefficient L_{ee} . Similarly, the thermal conductivity, κ , is given by [25]

$$\kappa = \frac{L_{QQ}L_{ee} - L_{Qe}L_{eQ}}{L_{ee}},\tag{21}$$

where the Onsager coefficients are given in Appendix and shown in Fig. 4.

The thermopower α is generally defined as the ratio of the *z* component of the electric field to the negative of the thermal gradient (also assumed to be in the *z* direction) under the condition of zero electrical current in the *z* direction. We write this condition as $\mathcal{E} = \alpha(-\nabla T)_{\mathbf{J}_{e}=0}$ [25]. It is readily shown that the α can be expressed in terms of the Onsager coefficients as

$$\alpha = -\frac{L_{eQ}}{L_{ee}}.$$
(22)

We can also calculate the coefficient L_{SQ} , which describes spin current parallel to the magnetic field and driven by a temperature gradient. A possible geometry for measuring this coefficient is shown in Fig. 1, as already mentioned. In this geometry, the *n*-type InSb is placed between two nonmagnetic contacts in an open circuit (no charge current), and a temperature difference is maintained between the two contacts.

The analysis of spin and thermal transport in this configuration has been discussed in Ref. [23] for a ferromagnetic metallic wire, and can also be used for the present case. As discussed there, the results depend on the boundary conditions at the ends of the wire, which are likely to fall between two possible limiting cases. In our notation, these are (i) $\Delta \mu = 0$ at each end of the wire; and (ii) $\mathbf{J}_S = 0$ at the two ends. For case (i), we can solve Eqs. (3) taking $\mathbf{J}_e = 0$ and $\Delta \mu = 0$ to obtain $\mathcal{E} = (L_{eQ}/L_{ee})\nabla T$ and

$$\mathbf{J}_{S} = \left(L_{SQ} - \frac{L_{Se}L_{eQ}}{L_{ee}}\right)(-\nabla T).$$
(23)

In our notation, this is equivalent to the result obtained in Ref. [23] for position-independent temperature gradient and boundary condition (i). For boundary condition (ii), one can solve Eq. (3) under the conditions $J_e = 0$, $J_S = 0$ to obtain

$$\mathcal{E} = \frac{L_{SS}L_{eQ} - L_{eS}L_{SQ}}{L_{ee}L_{SS} - L_{eS}L_{Se}}$$
(24)

and

$$-\frac{\hbar}{2q}\nabla(\Delta\mu) = \frac{L_{ee}L_{SQ} - L_{Se}L_{eQ}}{L_{ee}L_{SS} - L_{eS}L_{Se}}.$$
(25)

Thus, various observable quantities (spin current density, induced electric field) are sensitive to the spin-dependent Seebeck coefficient L_{SQ} . The results would be modified when spin-flip scattering is included, as described in Ref. [23]. The thermally induced spin accumulation (i.e., $\Delta \mu$) or spin current can also be detected nonlocally, by allowing the spin to diffuse into a nonmagnetic material, as discussed by Slachter *et al.* [24].

TABLE I. Numerical parameters used in the calculation of the Onsager coefficients given in Appendix. The estimate $\tau \sim 10^{-12}$ s is inferred from the electron mobility quoted in Ref. [1].

Assumed values for the physical properties of InSb		
Quantity	Value	Ref.
g	-49.0	[27]
m^*	$0.013m_{e}$	[19]
τ	$\approx 1 \times 10^{-12}$ s	[1]

IV. NUMERICAL RESULTS

We now turn to numerical results based on the present simplified model. We first calculate the chemical potential $\mu(T,B)$ at T = 0 K, assuming parameters appropriate to the conduction band of InSb and the experiments of Ref. [1], as given in Table I.

The resulting Fermi energy is shown in Fig. 2 as a function of *B*, assuming a conduction electron density of $\rho = 3.7 \times 10^{15}$ cm⁻³, as used in the experiments of Ref. [1]. The results show, as already obtained in Ref. [1], that $\mu(T = 0, B)$ is a nonmonotonic function of *B*, with discontinuous changes in slope wherever the minimum of one of the spin subbands rises through the Fermi energy and becomes unoccupied.

Given the Fermi energy, or at finite temperature the chemical potential $\mu(T, B)$, we can calculate a variety of transport coefficients. Here we calculate the components of electrical and thermal conductivities, and of the thermopower, parallel to the field, under the appropriate experimental conditions as described above. We also calculate the transport coefficient L_{SQ} [Eq. (3)], which represents the spin current density in the z direction per unit applied temperature gradient in the z direction. Expressions for the relevant Onsager coefficients are given in Appendix. All the coefficients are functions of both the applied magnetic field B and the temperature T. The



FIG. 2. (Color online) Calculated Fermi energy at T = 0 K plotted as a function of an applied magnetic field along the z direction as indicated in Fig. 1. We also show the first three Landau bands (n = 0, 1, 2); each band is labeled by its band index n and spin σ_z as (n, σ_z) , where $\sigma_z = \pm$. Straight lines correspond to the minima of the various Landau subbands, as labeled in the figure. Scalloped curve represents the T = 0 Fermi energy E_F as a function of magnetic field. In this figure, E_F is calculated neglecting spin-orbit coupling.



FIG. 3. The calculated electrical conductivity σ_e , given in Eq. (A1), plotted as a function of the inverse magnetic field 1/B at T = 4.5 K, and neglecting spin-orbit interactions. The maxima in the conductivity occur when the Fermi energy crosses the bottom of the Landau band as plotted in Fig. 2. The highest-field maximum occurs at approximately B = 1.2 T, which corresponds to the Fermi energy crossing the (0, -) Landau level. This curve is calculated neglecting spin-orbit interaction and using the T = 0 Fermi energy.

integrals in the transport coefficients are all dominated by energies within $k_B T$ of E_F , since the energy derivative of the Fermi function, which is a factor in each of the integrals, is strongly peaked near E_F .

The results of these calculations are shown in Figs. 3-6. In each case, we have plotted the transport coefficients at T = 4.5 K as functions of the inverse magnetic field. The various numerical parameters used in the calculations are given in Table I. We plot the transport coefficients in this manner in order to show that the positions of the peaks in these quantities vary periodically with 1/B. The oscillations are examples of the de Haas- van Alphen-like oscillations normally seen in the magnetotransport coefficients of metals [28]. In particular, the spin-dependent Seebeck coefficient L_{SO} shows these oscillations, which are similar to those seen in experiments (but the experiments measure the transverse spin-dependent Seebeck coefficient).



FIG. 4. Calculated thermal conductivity, κ , given in Eq. (21), plotted as a function of 1/B at T = 4.5 K. The maxima in κ occur, like those of σ_e , when the field-dependent Fermi energy crosses the bottom of a Landau level. This curve is calculated assuming no spin-orbit interaction and the values of the T = 0 K Fermi energy.

 0.15×10^{-4} 0.1 α (V/m-K) 0.05 0.0 0 1 2 3 4 5 $1/B(T^{-1})$

PHYSICAL REVIEW B 90, 174435 (2014)

FIG. 5. The calculated thermopower, as obtained using Eq. (22), plotted versus 1/B at T = 4.5 K. The maxima occur in the same manner as in Figs. 3 and 4. This curve is calculated assuming no spin-orbit interaction and the values of the T = 0 K Fermi energy.

V. DISCUSSION AND CONCLUSIONS

In this paper, we have given a simple model for the spin-dependent Seebeck coefficient in InSb. In our model, the electronic energy levels of *n*-type InSb in a magnetic field are given as Landau levels and the various electronic transport coefficients, including the spin-dependent Seebeck coefficient, are obtained from a simple Boltzmann equation approach for each Landau subband. The oscillations of this coefficient in a magnetic field occur when the Fermi energy crosses the minima of the various Landau subbands as the magnetic field is varied.

While our model holds, in principle, for any temperature T, we have carried out the calculations of the transport coefficients only at low T (T ~ 4.5 K) and specifically calculated the chemical potential at T = 0 K (see Fig. 2). While the difference between the chemical potential at T = 0and T = 4.5 K is small, it could affect both the magnitude and position of the Landau level crossings. This change could give quantitatively different results, but the qualitatively picture of the oscillations would remain the same.



FIG. 6. The calculated spin-dependent Seebeck coefficient L_{SQ} given by Eq. (A4), plotted versus 1/B at T = 4.5 K but using the T = 0 K Fermi energy. The maxima occur when the minima of the various Landau subbands cross the Fermi energy, as in Figs. 3-5.

NICHOLAS A. PIKE AND DAVID STROUD

Finally, we discuss how our simple model might be modified to produce a transverse spin-dependent Seebeck effect. The present model omits spin-orbit interaction, which is known to have a large effect on the band structure of InSb and similar compound semiconductors. The spin-orbit interaction couples the spatial momentum to various components of the electronic spin. In particular, some forms of this interaction couple momenta in one direction with spin components in other directions. Such coupling could lead to expectation values of the spin vector which are tilted relative to the electronic momentum. This could, in turn, produce a nonzero value of $\mathbf{J}_{S,x}$ and $\mathbf{J}_{S,y}$ along the z direction. If the spin vector is tilted relative to the direction of spin current, this will lead to a transverse electric field via the inverse spin Hall effect (ISHE) [29]. An ISHE electric field would also be produced if the spin is oriented in the z direction but the corresponding spin current has a component in the x or y direction. A model for this transverse spin-dependent Seebeck effect based on this picture is a subject for future work.

ACKNOWLEDGMENTS

This work was supported by the Center for Emergent Materials, an NSF MRSEC under Award No. DMR-1420451. The authors would like to thank C. Jaworski, J. P. Heremans, E. Johnston-Halperin, and R. C. Myers for valuable discussions.

APPENDIX: ONSAGER COEFFICIENTS

Here we give expressions for the various Onsager coefficients discussed and calculated in the text. In our model, the Onsager coefficients are

$$L_{ee} = \frac{L_z}{V} \sum_{n,\sigma_z} \int_{-\infty}^{\infty} \frac{dk_z}{2\pi} e^2 N_{n,\sigma_z} \left[v_{n,\sigma_z}(k_z) \right]^2 \tau \frac{\partial g^0_{n,\sigma_z}(E,T)}{\partial E};$$
(A1)

$$L_{eQ} = \frac{L_z}{V} \sum_{n,\sigma_z} \int_{-\infty}^{\infty} \frac{dk_z}{2\pi} e N_{n,\sigma_z} [v_{n,\sigma_z}(k_z)]^2 \tau \frac{\Delta E}{T} \frac{\partial g_{n,\sigma_z}^0(E,T)}{\partial E};$$
(A2)

$$L_{Se,i} = -\frac{L_z}{V} \sum_{n,\sigma_z} \int_{-\infty}^{\infty} \frac{dk_z}{2\pi} e\mu_B \langle \sigma_i(k_z) \rangle N_{n,\sigma_z} \Big[v_{n,\sigma_z}(k_z) \Big]^2 \tau \frac{\partial g^0_{n,\sigma_z}(E,T)}{\partial E};$$
(A3)

$$L_{SQ,i} = -\frac{L_z}{V} \sum_{n,\sigma_z} \int_{-\infty}^{\infty} \frac{dk_z}{2\pi} \mu_B \langle \sigma_i(k_z) \rangle N_{n,\sigma_z} \Big[v_{n,\sigma_z}(k_z) \Big]^2 \tau \frac{\Delta E}{T} \frac{\partial g_{n,\sigma_z}^0}{\partial E};$$
(A4)

$$L_{QQ} = \frac{L_z}{V} \sum_{n,\sigma_z} \int_{-\infty}^{\infty} \frac{dk_z}{2\pi} N_{n,\sigma_z} \frac{(\Delta E)^2}{T} \left[v_{n,\sigma_z}(k_z) \right]^2 \tau \frac{\partial g_{n,\sigma_z}^0}{\partial E},\tag{A5}$$

where $L_{eQ} = L_{Qe}$ and $\Delta E = E_{n,\sigma_z}(k_z) - \mu$. In Eqs. (A1)–(A5), the derivative $\partial g^0_{n,\sigma_z}(E,T)/\partial E = -\beta e^{\beta(E-\mu)}/(e^{\beta(E-\mu)} + 1]^2$, with $\beta = 1/(k_BT)$, and $E = E_{n,\sigma_z}(k_z)$. In our calculations, the integrals given above are converted to integrals over energy using the relationship $dk_z = dE/[dE/dk_z]$. All the integrals are dominated

by the energy range within k_BT of the Fermi energy, because $\partial g^0_{n,\sigma_z}/\partial E$ is strongly peaked around $E = E_F$. In practice, the integrands all become vanishingly small beyond an energy of $\sim 3k_BT$ on either side of E_F . In all the above expressions, $v_{n,\sigma_z}(k_z)$ is obtained from Eq. (16).

- C. M. Jaworski, R. C. Myers, E. Johnston-Halperin, and J. P. Heremans, Nature (London) 487, 210 (2012).
- [2] K. Uchida, H. Adachi, T. Ota, H. Nakayama, S. Maekawa, and E. Saitoh, Appl. Phys. Lett. 97, 172505 (2010).
- [3] K. Uchida, S. Takahoshi, K. Harii, J. Ieda, W. Koshibae, K. Ando, S. Maekawa, and E. Saitoh, Nature (London) 455, 778 (2008).
- [4] G. E. W. Bauer, E. Saitoh, and B. J. van Wees, Nat. Mater. 11, 391 (2012).
- [5] Y. Takezoe, K. Hosono, A. Takeuchi, and G. Tatara, Phys. Rev. B 82, 094451 (2010).
- [6] B. Scharf, A. Matos-Abiague, I. Zutic, and J. Fabian, Phys. Rev. B 85, 085208 (2012).

- [7] I. Zutic, J. Fabian, and S. Das Sarma, Rev. Mod. Phys. 76, 323 (2004).
- [8] J. Fabian, A. Matos-Abiague, C. Ertler, P. Stano, and I. Zutic, Acta Phys. Slov. 57, 565 (2007).
- [9] M. Johnson and R. H. Silsbee, Phys. Rev. B 35, 4959 (1987).
- [10] M. Johnson and R. H. Silsbee, Phys. Rev. B 37, 5312 (1988).
- [11] M. Hatami, G. E. W. Bauer, Q. Zhang, and P. J. Kelly, Phys. Rev. B 79, 174426 (2009).
- [12] E. O. Kane, J. Phys. Chem. Sol. 1, 249 (1957).
- [13] G. Dresselhaus, Phys. Rev. 100, 580 (1955).
- [14] R. H. Parmenter, Phys. Rev. 100, 573 (1955).
- [15] L. M. Roth, S. H. Groves, and P. W. Wyatt, Phys. Rev. Lett. 19, 576 (1967).

- [16] L. M. Roth, Phys. Rev. 173, 755 (1968).
- [17] C. R. Pidgeon and S. H. Groves, Phys. Rev. 186, 824 (1969).
- [18] J. Alsmeier, Ch. Sikorski, and U. Merkt, Phys. Rev. B 37, 4314(R) (1988).
- [19] G. Dresselhaus, A. F. Kip, C. Kittel, and G. Wagoner, Phys. Rev. 98, 556 (1955).
- [20] S. D. Brechet and J.-P. Ansermet, Phys. Stat. Sol. RRL 5, 423 (2011).
- [21] K. Uchida, S. Takahashi, J. Ieda, K. Harii, K. Ikeda, W. Koshibae, S. Maekawa, and E. Saitoh, J. Appl. Phys. **105**, 07C908 (2009).
- [22] T. Valet and A. Fert, Phys. Rev. B 48, 7099 (1993).

- [23] M. Hatami, G. E. W. Bauer, S. Takahashi, and Ds. Maekawa, Sol. State Commun. 150, 480 (2010).
- [24] A. Slachter, F. L. Bakker, J. P. Adam, and B. J. van Wees, Nat. Phys. 6, 879 (2010).
- [25] N. W. Ashcroft and N. D. Mermin, *Solid State Physics* (Brooks/Cole, Belmont, California, 1976).
- [26] L. Onsager, Phys. Rev. 37, 405 (1931).
- [27] O. Madelung, Landolt-Börnstein, Numerical Data and Functional Relationships in Science and Technology, New Series Group III, Vol. 17, Subvol. A, Sec. 2.15 (Springer, Berlin, 1982).
- [28] D. Shoenberg, *Magnetic Oscillations in Metals* (Cambridge University Press, New York, 1984).
- [29] K. Ando and E. Saitoh, Nat. Commun. 3, 629 (2012).